





CONTAINS CONFIDENTIAL PATIENT INFORMATION Zinbryta (daclizumab)

Prior Authorization of Benefits (PAB) Form

Complete form in its entirety and fax to: Prior Authorization of Benefits Center at 1-844-512-9004 Provider Help Desk: 1-800-454-3730

1. PATIENT INFORMATION		2. PHYSICIAN INFORMATION		
Patient	name:	Prescribing physician:		
Patient ID #:		Physician address:		
Patient DOB:		Physician phone #:		
Date of Rx:		Physician fax #:		
Patient phone #:		Physician specialty:		
Patient email address:		Physician DEA:		
		Physician NPI #:		
		Physician email address:		
3. MEDICATION	4. STRENGTH	5. DIRECTIONS	6. QUANTITY PER 30 DAYS	
Zinbryta (daclizumab)			Specify:	
7. DIAGNOSIS:				
8. APPROVAL CRITERIA: CHECK ALL BOXES THAT APPLY				
NOTE: Any areas not filled out are considered not applicable to your patient and MAY AFFECT THE OUTCOME of this request.				
Prior authorization is required for daclizumab (Zinbryta). Payment will be considered if all of these conditions are met: 1) Patient has a diagnosis of a relapsing form of multiple sclerosis (MS)				
2) Patient is 18 years of age or older				
3) Patient has documentation of previous trials and therapy failures with two or more drugs indicated for MS treatment				
4) Patient does not have pre-existing hepatic disease or hepatic impairment (including hepatitis B or C)				
5) Baseline transaminases (ALT, AST) and bilirubin levels are obtained				
6) Patient does not have an ALT or AST at least 2 times the upper limit of normal (ULN)				
7) Patient does not have a history of autoimmune hepatitis or other autoimmune condition involving the liver				
8) Patient has been screened for TB and treated for TB if positive				
9) Daclizumab will be used as monotherapy				
10) Daclizumab will be dosed as 150 mg once monthly				
11) Prescriber, patient and pharmacy are enrolled in the Zinbryta REMS program				
12) The 72-hour emergency supply rule does not apply to daclizumab.				

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Lost or stolen medication replacement requests will not be authorized. If criteria for coverage are met, an initial

clinical response to daclizumab therapy is provided.

authorization will be given for 12 months. Additional authorizations will be considered when documentation of a positive

13)







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Patient name: Patient ID #:				
Document two or more previous treatment failures:				
Trial 1: Drug name and dose:	Trial dates:			
Reason for failure:				
Trial 2: Drug name and dose:	Trial dates:			
Reason for failure:				
Trial 3: Drug name and dose:	Trial dates:			
Reason for failure:				
Does patient have pre-existing hepatic disease or hepatic impairment? ☐ Yes ☐ No				
Have baseline transaminases (ALT, AST) been obtained? ☐ Yes (attach res	sults) 🗆 No			
Does patient have ALT or AST at least two times the upper limit of normal?	□ Yes □ No			
Does patient have a history of autoimmune hepatitis or other autoimmune condition involving the liver? ☐ Yes ☐ No Has patient been screened for TB and treated for TB if positive?				
☐ Yes, provide result and treatment if positive:	□ No			
Will daclizumab be used as monotherapy? ☐ Yes ☐ No				
Prescriber, patient and pharmacy are enrolled in the Zinbryta REMS Program:	∃Yes □ No			
Renewal requests: Provide documentation of positive clinical response to daclizumab therapy:				
Attach lab results and other documentation as necessary.				
9. PHYSICIAN SIGNATURE				
Prescriber or authorized signature	Date			
Prior authorization of benefits is not the practice of medicine or the substitute for the independent of treating physician can determine what medications are appropriate for a patient. Please regarding benefits, conditions, limitations and exclusions. The submitting provider certifies complete, and the requested services are medically indicated and necessary to the health of	efer to the applicable plan for the detailed information s that the information provided is true, accurate and			
Note: Payment is subject to member eligibility. Authorization does not guarantee payment.				
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